>AJ303077 ACCESSION:AJ303077 NID: gi 12666719 emb AJ303077.1 HSA303077 Homo sapiens mRNA for mitochondrial RNA splicing protein 3/4 (HMRS3/4 gene), 1448 bp cDNA splice variant Length = 1448

Score = 760 bits (1940), Expect = 0.0 Identities = 364/364 (100%), Positives = 364/364 (100%) Frame = +3

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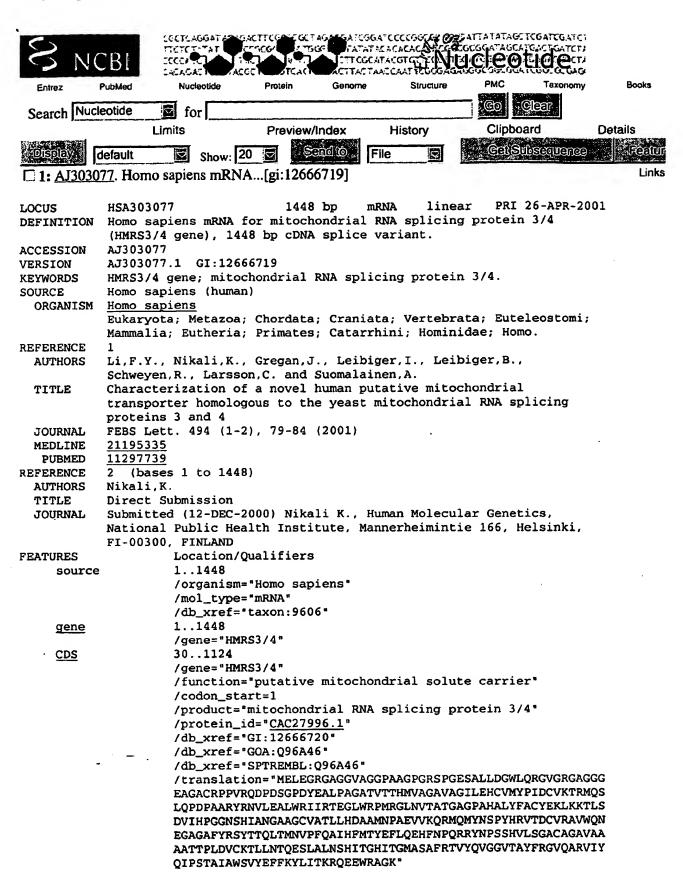
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Nuclei Sequence viewer rage 1 01 2



ORIGIN

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☐ 1: FEBS Lett. 2001 Apr 6;494(1-2):79-84.

Related Articles, Links

ELSEVIER FULL TEXT ARTICLE

Characterization of a novel human putative mitochondrial transporter homologous to the yeast mitochondrial RNA splicing proteins 3 and 4.

Li FY, Nikali K, Gregan J, Leibiger I, Leibiger B, Schweyen R, Larsson C, Suomalainen A.

Department of Molecular Medicine, CMM, Karolinska Hospital, Stockholm, Sweden. fangyuan.li@cmm.ki.se

We report here a novel human gene, hMRS3/4, encoding a putative mitochondrial transporter structurally and functionally homologous to the yeast mitochondrial RNA splicing proteins 3 and 4. These proteins belong to the family of mitochondrial carrier proteins (MCF) and are likely to function as solute carriers. hMRS3/4 spans approximately 10 kb of genomic DNA on chromosome 10q24 and consists of four exons that encode a 364-aa protein with six transmembrane domains. A putative splice variant, encoding a 177-aa protein with three transmembrane domains, was also identified. hMRS3/4 has a well-conserved signature sequence of MCF and is targeted into the mitochondria. When expressed in yeast, hMRS3/4 efficiently restores the mitochondrial functions in mrs3(o)mrs4(o) knock-out mutants. Ubiquitous expression in human tissues and a well-conserved structure and function suggest an important role for hMRS3/4 in human cells.

MeSH Terms:

- Alternative Splicing*
- Amino Acid Sequence
- Base Sequence
- Carrier Proteins/genetics*
- Carrier Proteins/metabolism
- Chromosome Mapping
- Chromosomes, Human, Pair 10*
- DNA, Complementary
- Fungal Proteins/genetics
- Gene Expression Profiling
- Human
- Membrane Transport Proteins*

- Mitochondria/metabolism*
- Molecular Sequence Data
- Mutagenesis
- Ophthalmoplegia, Chronic Progressive External/genetics
- RNA Splicing
- Repressor Proteins*
- Saccharomyces cerevisiae/genetics
- Saccharomyces cerevisiae Proteins*
- Spinocerebellar Ataxias/genetics
- Support, Non-U.S. Gov't
- Tissue Distribution
- Transfection

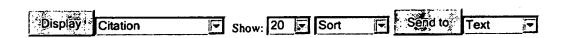
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- Carrier Proteins
- DNA, Complementary
- Fungal Proteins
- Membrane Transport Proteins
- Repressor Proteins
- SLC25A28 protein, human
- Saccharomyces cerevisiae Proteins
- MRS4 protein, S cerevisiae
- MRS3 protein, S cerevisiae

Secondary Source ID:

- GENBANK/AF327402
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- GENBANK/AJ303077
- GENBANK/AJ303078

PMID: 11297739 [PubMed - indexed for MEDLINE]



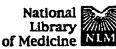
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□ 1: J Mol Biol. 1991 Jan 5;217(1):23-37.

Abstract

Related Articles, Links

Text

MRS3 and MRS4, two suppressors of mtRNA splicing defects in yeast, are new members of the mitochondrial carrier family.

Sort

Wiesenberger G, Link TA, von Ahsen U, Waldherr M, Schweyen RJ.

Institut fur Mikrobiologie und Genetik Universitat Wien, Austria.

When present in high copy number plasmids, the nuclear genes MRS3 and MRS4 from Saccharomyces cerevisiae can suppress the mitochondrial RNA splicing defects of several mit-intron mutations. Both genes code for closely related proteins of about Mr 32,000; they are 73% identical. Sequence comparisons indicate that MRS3 and MRS4 may be related to the family of mitochondrial carrier proteins. Support for this notion comes from a structural analysis of these proteins. Like the ADP/ATP carrier protein (AAC), the mitochondrial phosphate carrier protein (PiC) and the uncoupling protein (UCP), the two MRS proteins have a tripartite structure; each of the three repeats consists of two hydrophobic domains that are flanked by specific amino acid residues. The spacing of these specific residues is identical in all domains of all proteins of the family, whereas spacing between the hydrophobic domains is variable. Like the AAC protein, the MRS3 and MRS4 proteins are imported into mitochondria in vitro and without proteolytic cleavage of a presequence and they are located in the inner mitochondrial membrane. In vivo studies support this mitochondrial localization of the MRS proteins. Overexpression of the MRS3 and MRS4 proteins causes a temperature-dependent petite phenotype; this is consistent with a mitochondrial function of these proteins. Disruption of these genes affected neither mitochondrial functions nor cellular viability. Their products thus have no essential function for mitochondrial biogenesis or for whole yeast cells that could not be taken over by other gene products. The findings are discussed in relation to possible functions of the MRS proteins in mitochondrial solute translocation and RNA splicing.

PMID: 1703236 [PubMed - indexed for MEDLINE]

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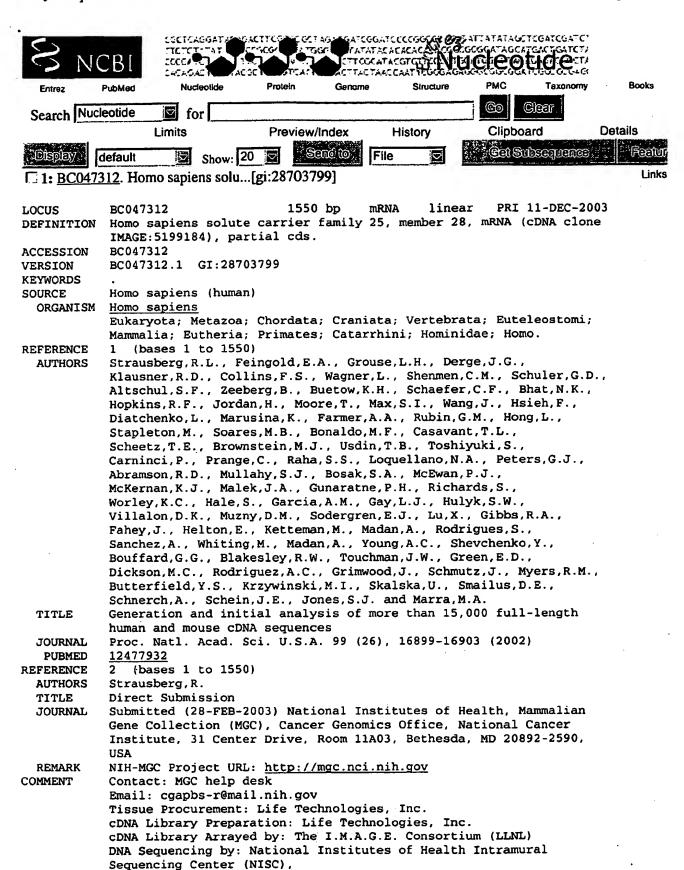
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Sbjct: 1157RAGK 1168

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            Contact: nisc_mgc@nhgri.nih.gov
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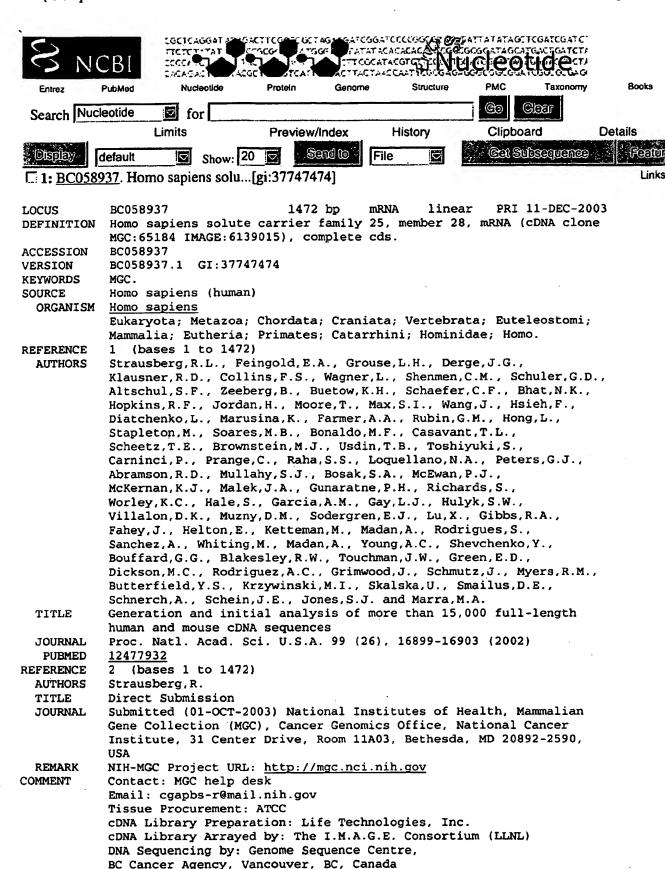
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Query: 361 RAGK 364

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Sbjct: 1119RAGK 1130

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rage Z OI 3

FEATURES

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CDS

ORIGIN

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info@bcgsc.bc.ca
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                /note="mito_carr; Region: Mitochondrial carrier protein"
                /db_xref="CDD:pfam00153"
                810..1115
misc_feature
                /gene="SLC25A28"
                /note=*mito_carr; Region: Mitochondrial carrier protein*
                /db_xref="CDD:pfam00153"
   1 gcggcggcgg gggtgggcgc cgcagctggc ccgggtggat ggagttggag gggcgggggtg
  61 ctggcggtgt ggcgggggg ccggcggcag ggcccgggcg gagccccggg gagtcggcgc
 121 tgctggacgg gtggctgcag cggggcgtgg gccggggggc cggcggcggg gaggccgggg
 181 cctgcaggcc cccggtacga caagatccgg actccggccc ggactacgag gcgctgccgg
 241 ctggagccac tgtcaccacg cacatggtgg caggcgccgt ggcagggatc ctggagcact
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301 gcgtgatgta ccccatcgac tgcgtcaaga cccggatgca gagtctacag cctgacccag

11

```
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481 attttgcctg ctacgaaaag ttaaaaaaga cattgagtga tgtaatccac cctgggggca
541 atagccatat tgccaatggt gcggccgggt gtgtggcaac attacttcat gatgcagcca
601 tgaaccctgc ggaagtggtc aagcagagga tgcagatgta caactcacca taccaccggg
661 tgacagactg tgtacgggca gtgtggcaaa atgaaggggc cggggccttt taccgcagct
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1021 tgcaggccag agtaatttac cagatcccct ccacagccat cgcatggtct gtgtatgagt
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1261 tggtgttttg actaacacca gttcctgcca acctctgttg ccaccacctt tccttccagg
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1441 taaaaagcct ttaaattaaa aaaaaaaaaa aa
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Jan 29 2004 15:38:25

characterize the protein. A starting material that can only be used to produce a final product does not have a substantial asserted utility in those instances where the final product is not supported by a specific and substantial utility. In this case none of the proteins that are to be produced as final products resulting from processes involving the claimed cDNA have asserted or identified specific and substantial utilities. The research contemplated by Applicants to characterize potential protein products, especially their biological activities, does not constitute a specific and substantial utility. Identifying and studying the properties of the protein itself or the mechanisms in which the protein is involved does not define a "real world" context of use. Note, because the claimed invention is not supported by a specific and substantial asserted utility for the reasons set forth above, credibility has not been assessed. Neither the specification as filed nor any art of record discloses or suggests any property or activity for the cDNA compounds such that another non-asserted utility would be well established for the compounds.

Claim 1 is also rejected under 35 U.S.C. § 112, first paragraph.

Specifically, since the claimed invention is not supported by either a specific and substantial asserted utility or a well established utility for the reasons set forth above, one skilled in the art would not know how to use the claimed invention.

Example 10: <u>DNA Fragment encoding a Full Open Reading Frame</u> (ORF)

Specification: The specification discloses that a cDNA library was prepared from human kidney epithelial cells and 5000 members of this library were

sequenced and open reading frames were identified. The specification discloses a Table that indicates that one member of the library having SEQ ID NO: 2 has a high level of homology to a DNA ligase. The specification teaches that this complete ORF (SEQ ID NO: 2) encodes SEQ ID NO: 3. An alignment of SEQ ID NO: 3 with known amino acid sequences of DNA ligases indicates that there is a high level of sequence conservation between the various known ligases. The overall level of sequence similarity between SEQ ID NO: 3 and the consensus sequence of the known DNA ligases that are presented in the specification reveals a similarity score of 95%. A search of the prior art confirms that SEQ ID NO: 2 has high homology to DNA Ligase encoding nucleic acids and that the next highest level of homology is to alpha-actin. However, the latter homology is only 50%. Based on the sequence homologies, the specification asserts that SEQ ID NO: 2 encodes a DNA ligase.

Claim 1: An isolated and purified nucleic acid comprising SEQ ID NO: 2.

Analysis: The following analysis includes the questions that need to be asked according to the guidelines and the answers to those questions based on the above facts:

1) Based on the record, is there a "well established utility" for the claimed invention? Based upon applicant's disclosure and the results of the PTO search, there is no reason to doubt the assertion that SEQ ID NO: 2 encodes a DNA ligase. Further, DNA ligases have a well-established use in the molecular biology art based on this class of protein's ability to ligate DNA. Consequently the answer to the question is yes.

Note that if there is a well-established utility already associated with the claimed invention, the utility need not be asserted in the specification as filed. In order to determine whether the claimed invention has a well-established utility the examiner must determine that the invention has a specific, substantial and credible utility that would have been readily apparent to one of skill in the art. In this case SEQ ID NO: 2 was shown to encode a DNA ligase that the artisan would have recognized as having a specific, substantial and credible utility based on its enzymatic activity.

Thus, the conclusion reached from this analysis is that a 35 U.S.C. § 101 rejection and a 35 U.S.C. § 112, first paragraph, utility rejection should not be made.

Example 11: Animals with Uncharacterized Human Genes

Specification: Kidney cells from a patient with Polycystic Kidney (PCK) Disease have been used to make a cDNA library. From this library 8000 nucleotide "fragments" have been sequenced but not yet used to express proteins in a transformed host cell nor have they been characterized in any other way. The 50 longest fragments, SEQ ID NO: 1-50, respectively, have been used to make transgenic mice. None of the 50 lines of mice have developed Polycystic Kidney Disease to date. The asserted utility is the use of the mice to research human genes from diseased human kidneys. The disease is inheritable, but chromosomal loci have not yet been identified. Neither the absence or presence of a specific protein has been identified with the disease condition.

App Serial # 09/870,113 Exhibit G Turner et at. Novel Human Mitochondriat Proteins and Polymucleotides Encoding the Sam

>AJ303077 ACCESSION:AJ303077 NID: gi 12666719 emb AJ303077.1 HSA303077 Homo sapiens mRNA for mitochondrial RNA splicing protein 3/4 (HMRS3/4 gene), 1448 bp cDNA splice variant Length = 1448

Score = 405 bits (1030), Expect = e-111 Identities = 192/192 (100%), Positives = 192/192 (100%) Frame = +3

- Query: 1 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 60 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG
- Sbjct: 30 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 209
- Query: 61 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 120 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR
- Sbjct: 210 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 389
- Query: 121 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 180 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA
- Sbjct: 390 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 569
- Query: 181 TLLHDAAMNPAE 192 TLLHDAAMNPAE Sbjct: 570 TLLHDAAMNPAE 605
- >BC047312 ACCESSION:BC047312 NID: gi 28703799 gb BC047312.1 Homo sapiens solute carrier family 25, member 28, mRNA (cDNA clone IMAGE:5199184), partial cds
 Length = 1550

Score = 405 bits (1030), Expect = e-111Identities = 192/192 (100%), Positives = 192/192 (100%) Frame = +2

- Query: 1 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 60 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG
- Sbjct: 77 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 256
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 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR
- Sbjct: 257 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 436
- Query: 121 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 180 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA
- Sbjct: 437 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 616

Query: 181 TLLHDAAMNPAE 192 TLLHDAAMNPAE

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>BC058937 ACCESSION:BC058937 NID: gi 37747474 gb BC058937.1 Homo sapiens solute carrier family 25, member 28, mRNA (cDNA clone MGC:65184 IMAGE:6139015), complete cds Length = 1472

Score = 405 bits (1030), Expect = e-111 Identities = 192/192 (100%), Positives = 192/192 (100%) Frame = +3

Query: 1 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 60 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG

Sbjct: 39 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 218

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Sbjct: 219 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 398

Query: 121 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 180

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Query: 181 TLLHDAAMNPAE 192

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Sbjct: 579 TLLHDAAMNPAE 614

>AJ303077 ACCESSION:AJ303077 NID: gi 12666719 emb AJ303077.1 HSA303077 Homo sapiens mRNA for mitochondrial RNA splicing protein 3/4 (HMRS3/4 gene), 1448 bp cDNA splice variant Length = 1448

Score = 405 bits (1030), Expect = e-111 Identities = 192/192 (100%), Positives = 192/192 (100%) Frame = +3

Query: 1 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 60 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG

Sbjct: 30 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 209

Query: 61 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 120 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR

Sbjct: 210 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 389

Query: 121 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 180 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA Sbjct: 390 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 569

Query: 181 TLLHDAAMNPAE 192 TLLHDAAMNPAE Sbjct: 570 TLLHDAAMNPAE 605

>BC047312 ACCESSION:BC047312 NID: gi 28703799 gb BC047312.1 Homo sapiens solute carrier family 25, member 28, mRNA (cDNA clone IMAGE:5199184), partial cds
Length = 1550

Score = 405 bits (1030), Expect = e-111 Identities = 192/192 (100%), Positives = 192/192 (100%) Frame = +2

Query: 1 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 60 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG

Sbjct: 77 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 256

Query: 61 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 120 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR Sbjct: 257 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 436

Query: 121 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 180 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA

Sbjct: 437 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 616

Query: 181 TLLHDAAMNPAE 192 TLLHDAAMNPAE Sbjct: 617 TLLHDAAMNPAE 652 >BC058937 ACCESSION: BC058937 NID: gi 37747474 gb BC058937.1 Homo sapiens solute carrier family 25, member 28, mRNA (cDNA clone MGC:65184 IMAGE:6139015), complete cds Length = 1472

Score = 405 bits (1030), Expect = e-111 Identities = 192/192 (100%), Positives = 192/192 (100%) Frame = +3

MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 60 Query: 1 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG

Sbjct: 39 MELEGRGAGGVAGGPAAGPGRSPGESALLDGWLQRGVGRGAGGGEAGACRPPVRQDPDSG 218

Query: 61 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 120 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR

Sbjct: 219 PDYEALPAGATVTTHMVAGAVAGILEHCVMYPIDCVKTRMQSLQPDPAARYRNVLEALWR 398

Query: 121 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 180

IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA

Sbjct: 399 IIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTLSDVIHPGGNSHIANGAAGCVA 578

Ouery: 181 TLLHDAAMNPAE 192

TLLHDAAMNPAE

Sbjct: 579 TLLHDAAMNPAE 614

>AJ303077 ACCESSION:AJ303077 NID: gi 12666719 emb AJ303077.1 HSA303077 Homo sapiens mRNA for mitochondrial RNA splicing protein 3/4 (HMRS3/4 gene), 1448 bp cDNA splice variant Length = 1448

Identities = 93/93 (100%), Positives = 93/93 (100%), Gaps = 0/93 (0%) Frame = +3

Query: 1 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL 60 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL 506

Query: 61 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE 93 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE Sbjct: 507 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE 605

>BC047312 ACCESSION:BC047312 NID: gi 28703799 gb BC047312.1 Homo sapiens solute carrier family 25, member 28, mRNA (cDNA clone IMAGE:5199184), partial cds
Length = 1550

Identities = 93/93 (100%), Positives = 93/93 (100%), Gaps = 0/93 (0%) Frame = +2

Query: 1 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL 60 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL Sbjct: 374 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL 553

Query: 61 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE 93 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE Sbjct: 554 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE 652

>BC058937 ACCESSION:BC058937 NID: gi 37747474 gb BC058937.1 Homo sapiens solute carrier family 25, member 28, mRNA (cDNA clone MGC:65184 IMAGE:6139015), complete cds Length = 1472

Identities = 93/93 (100%), Positives = 93/93 (100%), Gaps = 0/93 (0%) Frame = +3

Query: 1 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL 60 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL Sbjct: 336 MQSLQPDPAARYRNVLEALWRIIRTEGLWRPMRGLNVTATGAGPAHALYFACYEKLKKTL 515

Query: 61 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE 93 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE Sbjct: 516 SDVIHPGGNSHIANGAAGCVATLLHDAAMNPAE 614

Query= SEQ ID NO:1 (1095 letters)

Sequences producing significant alignments:	Score (bits)	E Value				
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Identities = 520/520 (100%) Strand = Plus / Minus						
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Query: 636 acgggcagtgtggcaaaatgaaggggccggggccttttaccgcagctacacca		•				
Query: 696 gaccatgaacgttcctttccaagccattcacttcatgacctatgaattcctgc						
Query: 756 ctttaacccccagagacggtacaacccaagctcccacgtcctctctggagctt	gcgcagg					
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Query: 876 ggagtccttggctttgaactcacacattacaggacatatcacaggcatggcta	gtgcctt					
		82631				
Query: 936 caggacggtatatcaagtaggtggggtgaccgcctatttccgaggggtgcagg						
Sbjct: 82630 caggacggtatatcaagtaggtgggtgaccgcctatttccgaggggtgcagg Query: 996 aatttaccagatccctccacagccatcgcatggtctgtgtatgagttcttca						
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Strand = Plus / Minus

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Sbjct: 91837 gccggcggcggggaggccggggcctgcaggccccggtacgacaagatccggactccggc 91778

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Sbjct: 91777 ccggactacgaggcgctgccggctggagccactgtcaccacgcacatggtggcaggcgcc 91718

Query: 241 gtggcagggatcctggagcactgcgtgatgtaccccatcgactgcgtcaag 291

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Identities = 233/233 (100%)

Strand = Plus / Minus

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Identities = 58/58 (100%) Strand = Plus / Minus

Query= SEQ ID NO:3 (582 letters)

Query: 350

Score E (bits) Value Sequences producing significant alignments: 577 e-162 AL353719.10.1.123160 >AL353719.10.1.123160 Length = 123160Score = 577 bits (291), Expect = e-162Identities = 291/291 (100%) Strand = Plus / Minus Query: 1 Query: 61 ${\tt gccggcggcggggaggccggggcctgcaggccccggtacgacaagatccggactccggc\ 180}$ Query: 121 Sbjct: 91837 gccggcggggggggggggcctgcaggccccggtacgacaagatccggactccggc 91778 ccggactacgaggcgctgccggctggagccactgtcaccacgcacatggtggcaggcgcc 240 Query: 181 Sbjct: 91777 ccggactacgaggcgctgccggctggagccactgtcaccacgcacatggtggcaggcgcc 91718 gtggcagggatcctggagcactgcgtgatgtaccccatcgactgcgtcaag 291 Query: 241 Sbjct: 91717 gtggcagggatcctggagcactgcgtgatgtaccccatcgactgcgtcaag 91667 Score = 462 bits (233), Expect = e-127Identities = 233/233 (100%) Strand = Plus / Minus agacccggatgcagagtctacagcctgacccagctgcccgctatcgcaatgtgttggagg 349 Query: 290 Sbjct: 85548 agacccggatgcagagtctacagcctgacccagctgcccgctatcgcaatgtgttggagg 85489

ccctctggaggattataagaacggagggcctatggaggcccatgagggggctgaacgtca 409

Sbjct: 85488 ccctctggaggattataagaacggagggcctatggaggcccatgagggggctgaacgtca 85429

cagcaacaggcgcagggcctgcccacgccctttattttgcctgctacgaaaagttaaaaa 469 Query: 410

Sbjct: 85428 cagcaacaggcgcagggcctgcccacgccctttattttgcctgctacgaaaagttaaaaa 85369

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Sbjct: 85368 agacattgagtgatgtaatccaccctgggggcaatagccatattgccaatggt 85316

Score = 117 bits (59), Expect = 8e-24

Identities = 59/59 (100%)

Strand = Plus / Minus

ggtgcggccgggtgtgtggcaacattacttcatgatgcagccatgaaccctgcggaagg 578 Query: 520

Sbjct: 84222 ggtgcggccgggtgtgtggcaacattacttcatgatgcagccatgaaccctgcggaagg 84164

Query= SEQ ID NO:5 (693 letters)

Query: 350

Score E (bits) Value Sequences producing significant alignments: 577 e-162 AL353719.10.1.123160 >AL353719.10.1.123160 Length = 123160Score = 577 bits (291), Expect = e-162Identities = 291/291 (100%) Strand = Plus / Minus atqqaqttqqaqqqcqqgqtgctggcggtgtggcggggggggccggcggcagggccggg 60 Query: 1 Ouery: 61 cqqaqcccqqqqaqtcgqcgctgctggacgggtggctgcagcgggggcgtgggggg 120 Sbjct: 91897 cggagccccggggagtcggcgctgctggacgggtggctgcagcgggggcgtgggccggggg 91838 gccggcggcggggaggccggggcctgcaggccccggtacgacaagatccggactccggc 180 Query: 121 Sbjct: 91837 gccggcggcgggggggccggggcctgcaggcccccggtacgacaagatccggactccggc 91778 $\verb|ccggactacgaggcgctgccggctggagccactgtcaccacgcacatggtggcaggcgcc|| 240$ Query: 181 Sbjct: 91777 ccggactacgaggcgctgccggctggagccactgtcaccacgcacatggtggcaggcgcc 91718 gtggcagggatcctggagcactgcgtgatgtaccccatcgactgcgtcaag 291 Query: 241 Sbjct: 91717 gtggcagggatcctggagcactgcgtgatgtaccccatcgactgcgtcaag 91667 Score = 462 bits (233), Expect = e-127Identities = 233/233 (100%) Strand = Plus / Minus agacceggatgcagagtctacagcctgacccagctgcccgctatcgcaatgtgttggagg 349 Query: 290

Sbjct: 85548 agacccggatgcagagtctacagcctgacccagctgcccgctatcgcaatgtgtttggagg 85489

Sbjct: 85488 ccctctggaggattataagaacggagggcctatggaggcccatgaggggctgaacgtca 85429

ccctctggaggattataagaacggagggcctatggaggcccatgagggggctgaacgtca 409

Query: 410 cagcaacaggcgcagggcctgcccacgcctttattttgcctgctacgaaaagttaaaaa 469

Sbjct: 85428 cagcaacaggcgcagggcctgcccacgcctttattttgcctgctacgaaaagttaaaaa 85369

Query: 470 agacattgagtgatgtaatccaccctgggggcaatagccatattgccaatggt 522

Sbjct: 85368 agacattgagtgatgtaatccaccctgggggcaatagccatattgccaatggt 85316

Score = 345 bits (174), Expect = 2e-92

Identities = 174/174 (100%)

Strand = Plus / Minus

Query: 520 ggtgcggccgggtgtgtggcaacattacttcatgatgcagccatgaaccctgcggaaggt 579

Sbjct: 84222 ggtgcggccgggtgtgtggcaacattacttcatgatgcagccatgaaccctgcggaaggt 84163

Query: 580 aatgattcctcaacctatcactctgtgggcagctgcacctgtatttctttacagtttgca 639

Sbjct: 84162 aatgattcctcaacctatcactctgtgggcagctgcacctgtatttctttacagtttgca 84103

Query: 640 gaagaaagcacatcagttttggtgggaaattctgttaccttgttttaccactag 693

ι

Sbjct: 84102 gaagaaagcacatcagttttggtgggaaattctgttaccttgtttaccactag 84049

Query= SEQ ID NO:11 (396 letters)

Sequences producing significant alignments:

AL353719.10.1.123160

Score E (bits) Value

>AL353719.10.1.123160 Length = 123160

Score = 446 bits (225), Expect = e-123 Identities = 225/225 (100%)
Strand = Plus / Minus

Query: 61 aggattataagaacggagggcctatggaggcccatgagggggctgaacgtcacagcaaca 120

Sbjct: 85480 aggattataagaacggagggcctatggaggcccatgagggggctgaacgtcacagcaaca 85421

Query: 121 ggcgcagggcctgcccacgccctttattttgcctgctacgaaaagttaaaaaagacattg 180

Sbjct: 85420 ggcgcagggcctgcccacgccctttattttgcctgctacgaaaagttaaaaaagacattg 85361

Query: 181 agtgatgtaatccaccctgggggcaatagccatattgccaatggt 225

Sbjct: 85360 agtgatgtaatccacctgggggcaatagccatattgccaatggt 85316

Score = 345 bits (174), Expect = 1e-92 Identities = 174/174 (100%) Strand = Plus / Minus

Query: 283 aatgattcctcaacctatcactctgtgggcagctgcacctgtatttctttacagtttgca 342

Query: 343 gaagaaagcacatcagttttggtgggaaattctgttaccttgttttaccactag 396

rage 1 of 50

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COMMENT						gi: 14280413.	200	
		During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations						
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	was ge	was generated from part of bacterial clone contigs of human						
		chromosome 10, constructed by the Sanger Centre Chromosome 10 Mapping Group. Further information can be found at						
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		35A1 It may bons only once				e overlapping		
	The tr	rue left end	of clone	RP11-85A1	is at 1 in	this sequence.	The	

true left end of clone RP11-483F11 is at 123061 in this sequence.